

### Background

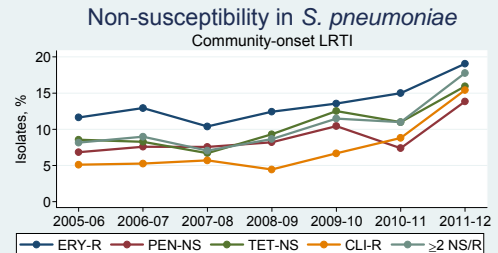
- The BSAC Respiratory Resistance Surveillance Programme has monitored antimicrobial susceptibility in the UK and the Republic of Ireland in the major pathogens causing community-onset and hospital-onset lower respiratory tract infections (LRTIs) since 1999/2000 and 2008/09, respectively.
- In 2011/12, the Programme collected 2,379 LRTI isolates from 38 clinical laboratories across the UK and Ireland.

### Methods

- Clinical laboratories collect up to a defined quota of isolates (excluding duplicates and isolates from cystic fibrosis patients) in each season (now Oct-Sept; was Oct-April up to 2007/08).
- MICs are measured and interpreted by BSAC methods.
- Hospital-onset isolates are those first obtained after >48 hours of hospital admission; all others are considered community-onset.
- See [www.bsacsurv.org](http://www.bsacsurv.org) or *JAC*, 2008, **62**, suppl 2 ii15 - ii28.

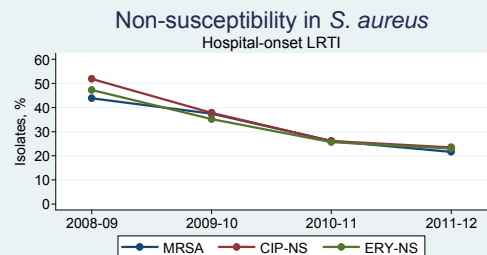
### 1,108 COMMUNITY-ONSET lower respiratory tract isolates collected October 2011 - Sept 2012 (in community, or hospital ≤48 hours)

- In 2011/12, 53 of 383 *S. pneumoniae* (14%) were non-susceptible (NS) to penicillin (all intermediate with MICs ≤2 mg/L). Seventy-four (19%) were NS to erythromycin, 64 being highly resistant with MICs ≥16 mg/L, and 59 (15%) were NS to clindamycin, continuing an apparent upward trend - see graph.
- As in earlier years, 21% of 489 *H. influenzae* and 99% of 236 *M. catarrhalis* produced β-lactamase, but remained very broadly (>95%) susceptible to other established antibiotic classes.



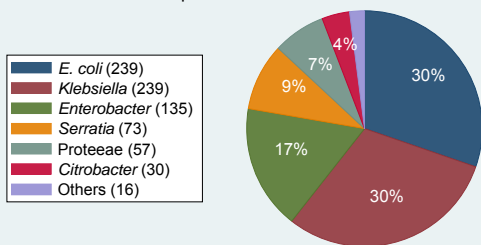
### 1,271 HOSPITAL-ONSET lower respiratory tract isolates collected October 2011 - Sept 2012 (>48 hours after admission to hospital)

- MRSA fell marginally in the last year to 22% of 209 *S. aureus* in 2011/12 after a rapid fall from 44% in 2008/09 to 26% in 2010/11.
- Non-susceptibility to ciprofloxacin and erythromycin fell with similar pattern to MRSA, reaching 23% and 23%, respectively.
- All *S. aureus* were susceptible to vancomycin, teicoplanin and linezolid in 2011/12.
- 38% of hospital-onset *S. aureus* LRTI were from intensive care units (ICU), similar to previous years, and MRSA was much less prevalent in ICU than other wards (10% vs 28%).



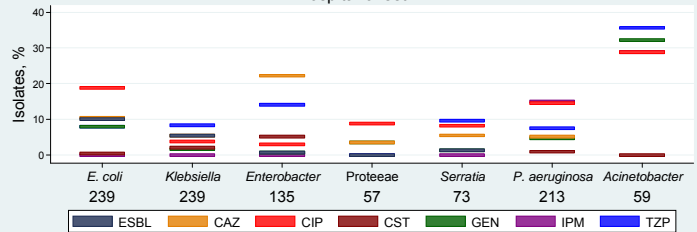
### Enterobacteriaceae, 2011/12

Hospital-onset LRTI: N = 789



### Non-susceptibility in Gram-negative bacteria, 2011/12

Hospital-onset LRTI



- In 2011/12, ESBLs were found in 10% of *E. coli*, 5% of *Klebsiella* and <1% of *Enterobacter* - rates similar to or lower than 2010/11 (11%, 11% and 5% respectively).
- Ciprofloxacin NS was also no higher than in 2010/11 (*E. coli* 19% vs 27%, *Klebsiella* 4% vs 7%, and *Enterobacter* 3% vs 7%).
- All isolates of **Enterobacteriaceae** from LRTI in 2011/12 were susceptible to imipenem (excluding 57 *Proteaeae* and 16 'Others').

- 5% of *Enterobacter* were **colistin**-non-susceptible, but only 2% of *Klebsiella* and <1% of *E. coli*, *Pseudomonas* and *Acinetobacter*.
- Non-susceptibility among 213 *P. aeruginosa* ranged up to 15% (imipenem and ciprofloxacin), similar to previous years.
- About 30% of 59 *Acinetobacter* were non-susceptible to imipenem, ciprofloxacin, gentamicin and piperacillin-tazobactam.

### CONCLUSIONS

- H. influenzae* and *M. catarrhalis* from community-onset LRTI remain very widely susceptible to existing antimicrobials.
- Community-onset *S. pneumoniae* also remain widely susceptible, despite a possible upward trend in resistance since about 2008.
- Non-susceptibility in isolates from hospital-onset LRTI in 2011/12 was generally similar to or lower than the three years before.

**Abbreviations:** NS non-susceptible (intermediate & resistant), R resistant. CAZ ceftazidime, CIP ciprofloxacin, CLI clindamycin, CST colistin, ERY erythromycin, GEN gentamicin, IPM imipenem, PEN penicillin, TET tetracycline, TZP piperacillin/tazobactam. ESBL extended-spectrum β-lactamase. MRSA methicillin-resistant *S. aureus*. LRTI lower respiratory tract infection.

**Extended Working Party Members (October 2013):** A. MacGowan<sup>1</sup> (Chair), M. Allen<sup>2</sup>, D. Brown<sup>3</sup>, P. Fernandes<sup>4</sup>, H. Grundmann<sup>5</sup>, R. Janes<sup>6</sup>, A. Johnson<sup>7</sup>, M. Jones<sup>8</sup>, A. Kidney<sup>6</sup>, D. Livermore<sup>7</sup>, S. McCurdy<sup>9</sup>, V. Martin<sup>1</sup>, T. Mephams<sup>10</sup>, S. Mushtaq<sup>7</sup>, S. Peacock<sup>11</sup>, J. Porter<sup>12</sup>, R. Reynolds<sup>1</sup>, C. Thomson<sup>13</sup>.

**Organism ID and Susceptibility Testing:** A. Kidney<sup>6</sup> and staff at Quotient BioAnalytical Sciences

<sup>1</sup>North Bristol NHS Trust; <sup>2</sup>Novartis; <sup>3</sup>EUCAST Scientific Secretary; <sup>4</sup>Cempra; <sup>5</sup>RIVM; <sup>6</sup>Quotient BioAnalytical Sciences, Fordham; <sup>7</sup>Public Health England, London; <sup>8</sup>Basilea; <sup>9</sup>Cubist; <sup>10</sup>AstraZeneca; <sup>11</sup>University of Cambridge; <sup>12</sup>Pfizer; <sup>13</sup>Astellas.

**Collecting Laboratories:** See [www.bsacsurv.org](http://www.bsacsurv.org) **Sponsors 2011/12:** Basilea, Cempra, Cubist, Pfizer. **Support:** BSAC